Supplementary material

Progesterone increases blood glucose via hepatic progesterone receptor membrane component 1 under limited or impaired action of insulin

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Fig S1. mRNA expression of *PGRMC1* and *PEPCK* in Hep3B cells AG-205 (0.1, 1, 10 μ g/ml) was treated in low glucose medium (50mg/dl) for 18 h in Hep3B cells. *RPLP0* was used as an internal control. Values represent means \pm SD. *, p<0.05 vs. Control. All experiments were repeated at least 3 times.



Fig S2. Western blot analysis and quantification of PGRMC1, phosphor/total CREB, and PEPCK in control and *PGRMC1* siRNA groups. Cells were starved in low glucose medium (50 mg/dl). P4 (10 nM) was treated for 18 h after siRNA transfection. β -actin was used as an internal control. Values represent means \pm SD. *, p<0.05 vs. Control. All experiments were repeated at least 3 times.



Fig S3. mRNA expression of *Pgrmc1* and *Pepck* in WT and *Pgrmc1* KO hepatocyte. Cells were incubated in low glucose medium (50mg/dl) for 18 h in Hep3B cells. *Rplp0* was used as an internal control. Values represent means \pm SD. *, p<0.05 vs. WT hepatocyte. All experiments were repeated at least 3 times.



Fig S4. Western blot analysis and quantification of PGRMC1 and PEPCK in Control vs. Dexamethasone, E2, P4, AG-205 groups. Cells were starved in DMEM medium (w/o glucose, w/o FBS). Dexamethasone (100 nM), E2 (100 nM), P4 (100 nM) and AG-205 (10 μ g/ml) were treated in DMEM medium (w/o glucose, 2%CD-FBS, 1% Penicillin/Streptomycin, 1nM dexamethasone) for 24 h. β -actin was used as an internal control. Values represent means \pm SD. *, p<0.05 vs. Control. All experiments were repeated at least 3 times.



















Phosphor CREB



Fig.4B	PGRMC1	РЕРСК	β-actin

















Fig.6C	PGRMC1	PEPCK	β-actin
	CREB	Phosphor CREB	

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PEPCK



β-actin



Fig.S2

Fig.S4	PGRMC1	PEPCK	β-actin